

XP-002170584

AN - 1988-074331 [11]

AP - JP19860171364 19860721

CPY - ASAH

DC - B04 D16

FS - CPI

IC - C12N5/00 ; C12N15/00 ; C12P21/02 ; C12R1/91

MC - B04-B04A D05-H08 D05-H12

M1 - [01] M423 M720 M903 N135 N136 N513 Q233 V753 V754; 3102-R

PA - (ASAH) ASAHI CHEM IND CO LTD

PN - JP63028386 A 19880206 DW198811 010pp

PR - JP19860171364 19860721

XA - C1988-033353

XIC - C12N-005/00 ; C12N-015/00 ; C12P-021/02 ; C12R-001/91

AB - J63028386 A transforming growth factor beta gene combined to a promoter is introduced in an animal cell to culture the animal cell and the resultant culture growth cell is used.

- USE/ADVANTAGE - The method can prepare the useful substance commercially.

- In an example, Human TGF beta gene was prepd. from human nasopharynx cancer-originated KB cell (ATCC CCL-17). Plasmid pTS neo was prepd. by process in which a terminator of T-antigen gene of SV 40 was connected to neomycin-resistant gene. Plasmid pMGSneo was prepd. by a process in which mouse metallothionein gene promoter was connected to TFG beta. Plasmid pMGRneo was prepd. by a process in which TGF beta gene was connected to the downstream of LTR of RSV. NIH3T3H cell was transformed by DNA of Plasmid pMGSneo and 2 clones were selected (NMTG 1 and NMTG 2). Number of cell after 4 days culture at 37 deg.C were as follows: (Clone, Number of cell/ml): NIH3T3 5.8 x 10 power 6; NMTG1 3.3 x 10 power 7; NMTG2 1.8 x 10 power 7.(0/3)

DRL - 3102-R

IW - CELL CULTURE PROCESS INTRODUCING TRANSFORM GROWTH FACTOR BETA GENE COMBINATION PROMOTE ANIMAL CELL

IKW - CELL CULTURE PROCESS INTRODUCING TRANSFORM GROWTH FACTOR BETA GENE COMBINATION PROMOTE ANIMAL CELL

NC - 001

OPD - 1986-07-21

ORD - 1988-02-06

PAW - (ASAH) ASAHI CHEM IND CO LTD

TI - Cell culture process - involves introduction of transforming growth factor beta gene combining to promoter into animal cell